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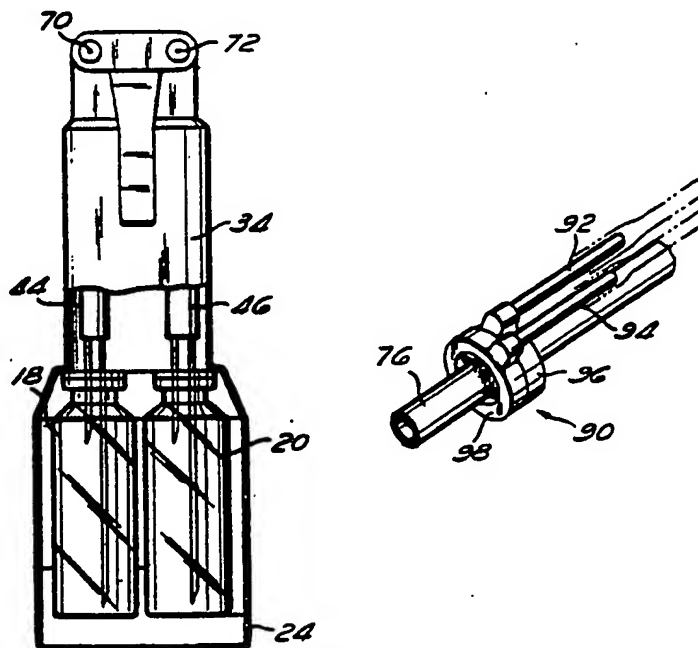
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(54) Title: FIBRINOGEN DISPENSING KIT



(57) Abstract

Disclosed is a dispenser (10) for separately dispensing each of two biological fluids contained therein for intermixing at a side outside of the dispenser to produce hemostasis or a tissue adhesive. The dispenser is compact, contains internal reservoirs (18, 20) of the two biological fluids, and is designed for efficient reloading with filled nonsterile containers without comprising the integrity of the sterile field. The dispenser is capable of dispensing the biological fluids, such as fibrinogen and thrombin, at either a focused point or in an aerosol mist. In addition, spray elements (82, 90, 96) are disclosed for uniformly distributing the two biological fluids along either the interior surface or the exterior surface of an implantable vascular graft.

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FIBRINOGEN DISPENSING KIT

Background of the Invention

5 The in vivo mechanism for the clotting of blood requires conversion of the soluble protein fibrinogen into the insoluble protein fibrin in a reaction catalyzed by the enzyme thrombin. Thrombin is the activated form of prothrombin, a globulin circulating in the plasma. The
10 conversion of prothrombin to thrombin requires a number of reactions involving the interaction of blood fractions having thromboplastic activity, including Stewart-Prower factor, Factor V, Factor VII and calcium. Several additional substances having thromboplastic activity have
15 been identified, and are generally referred to by the genus thromboplastin.

 Under the influence of thrombin, which is actually a proteolytic enzyme, and other blood factors, the blood protein fibrinogen loses several polypeptide chains to form
20 fibrin. The fibrin then undergoes a polymerization to produce fibrin polymers which contribute to the physical properties of the clot.

 The thrombin catalyzed polymerization of fibrinogen into fibrin is reproducible in vitro. For example, a system
25 composed of purified fibrinogen, thrombin and added calcium produces the so-called Fibrin S polymer by spontaneous reversible polymerization. Polymerization is said to be reversible, due to the solubility of the Fibrin S polymer in dilute (0.03%) HCl.

30 However, an acid insoluble polymer of Fibrin I (insoluble fibrin) can be formed in vitro for example by addition of small amounts of serum to the in vitro system.

 The serum contains a factor responsible for inhibition of the reversible polymerization of fibrin. This
35 stabilizing factor in serum which enters the clotting sequence after fibrin has been formed, known as the Laki-Lorand factor (LLF) or fibrin stabilizing factor (FSF), is

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known to exhibit a useful stabilizing effect on the in vitro polymerization of fibrinogen. Under the nomenclature proposed by the International Committee for the Standardization of the Nomenclature of Blood Clotting Factors, LLF is designated Factor XIII.

Thus, it is known that purified thrombin and fibrinogen, together with a variety of known adjuvants, can be combined in vitro to produce a polymer having great potential benefit, both as a hemostatic agent and as a tissue adhesive. Because of the rapid polymerization upon intimate interaction of fibrinogen and thrombin, it is important to maintain these two blood proteins separate until the application site. Previously, these materials have been delivered by devices such as a dual syringe apparatus which makes it possible to deliver the fluids for mixture at a small point.

One apparatus for applying a fibrinogen-based tissue adhesive is disclosed in U.S. Patent No. 4,359,049 to Redl, et al. Redl discloses a complicated mechanism in which two standardized one-way syringes are held in a support having a common actuating means guided along a rod. Each of the delivery ends of the syringes is inserted into a collection manifold for delivery of the two components for mixing and delivery optimally through a mixing needle. However, it is often desirable or necessary to cover a broad area of a wound, either to stop bleeding, to fix tissue or to prevent infection.

Notwithstanding the contribution of Redl, there remains a need for a fibrinogen dispensing kit which is inexpensive, suitable for entry into the sterile field, and adaptable for convenient recharging of the fibrinogen or thrombin from a single source, as may be required during prolonged surgery or other procedure. In addition, the dispensing kit is advantageously capable of delivering the two blood proteins either at a focused point of delivery or as an aerosol mist for covering a region of tissue.

Summary of the Invention

There has been provided in accordance with one aspect of the present invention a dispenser for separately dispensing each of two biological fluids contained therein, for intermixing at a remote site to produce hemostasis or a tissue adhesive, depending upon the application. The dispenser comprises a housing, having a first and a second pump therein. A first hollow aspirating needle in the housing is in fluid communication with the first pump, and a second hollow aspirating needle is in fluid communication with the second pump. The first and second needles extend within the housing into a chamber adapted for removably receiving a first and a second reservoir of biological fluids to be dispensed.

In a preferred embodiment, each of the first and second aspirating needles terminates in a sharp point, for piercing a septum on the reservoir of biological fluid. Preferably, one or both of the reservoirs comprise a standard vacuum bottle having a piercable, resealable septum as a closure member. The dispenser further comprises a first and a second hollow vent needle in the housing, extending into the chamber substantially parallel to the first and second aspirating needles, respectively, for communicating with the interior of the reservoirs.

In one embodiment of the present invention, the housing is further provided with a common actuator for simultaneously actuating each of the first and second pumps. The actuator is independently removably linked to each pump, to permit partial disengagement thereof for independent operation of the pumps. A cover is provided on the housing, for enclosing the chamber for receiving the reservoirs of biological fluid. The cover may be pivotally attached to the housing, or may be entirely removable from the housing. The cover comprises a cavity adapted to receive at least one of the reservoirs so that the cover

may be loaded with the reservoirs and then installed as a unit into the housing.

In accordance with another aspect of the present invention, there is provided an interface device for
5 placing a vacuum bottle having a resealable septum thereon in valved fluid communication with a source of fluid. The interface comprises a housing having a flow channel therethrough, and a valve in the flow channel. A hollow
10 needle is provided in communication with the flow channel on the first side of the valve, and a hollow cannulae is provided in communication with the channel on the second side of the valve. The needle is adapted for piercing the septum on a vacuum bottle.

In accordance with a further embodiment of the present
15 invention, there is provided a radial spray head for dispensing each of a plurality of fluids onto the interior surface of a tubular sleeve, comprising a first spray element having an influent passageway in fluid
20 communication with each of a plurality of effluent ducts disposed about the periphery of the first spray element, and a second spray element comprising a second influent duct in fluid communication with each of a plurality of effluent ducts disposed about the periphery of the second
25 spray element. The effluent ducts are disposed about the exterior periphery of the first and second spray elements for directing a spray in a radially outward direction.

For dispensing fluids on the exterior surface of a tubular sleeve, the first and second spray elements are provided with a cylindrical passageway therethrough for
30 receiving the tubular sleeve. Effluent ducts are disposed about the surface of the cylindrical passageway for directing fluid in a radially inward direction.

These and other features and advantages of the present invention will become apparent from the detailed
35 description of preferred embodiments which follows, when taken together with the attached drawings and claims.

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Brief Description of the Drawings

Figure 1 is an elevational perspective view of a dispenser according to one embodiment of the present invention.

5 Figure 2 is a perspective view of the embodiment illustrated in Figure 1, showing the cover open and exposing the container receiving areas.

10 Figure 3 is a partial sectional front elevational view of a dispenser according to a second embodiment of the present invention.

Figure 4 is a perspective view of the embodiment illustrated in Figure 3.

15 Figure 5 is a perspective view of an interface for loading the containers used in the dispenser of the present invention.

Figure 6 is a perspective view of a dual fluid radial dispenser according to the present invention, for dispensing fluid on the interior of a tubular vascular graft.

20 Figure 7 is a perspective view of a single element from the radial dispenser shown in Figure 6.

Figure 8 is a partial cross-sectional view of the dispenser shown in Figure 6.

25 Figure 9 is a radial dispenser for dispensing fluids on the exterior surface of a tubular sleeve.

Figure 10 is a perspective view of a single fluid dispensing element from the dispenser shown in Figure 9.

30 Figure 11 illustrates a spray pattern of a dispensing head according to the present invention, wherein intermixing occurs at a first distance from the spray head.

Figure 12 is a dispenser head according to the present invention, wherein intermixing occurs at a second distance from the spray head.

35 Figure 13 illustrates a third embodiment of the spray head, wherein intermixing occurs at a focused point.

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Figure 14 illustrates a rigid attachment for conducting blood proteins to a remote location.

Figure 15 illustrates an elongate flexible attachment for conducting blood proteins from the spray head of the present invention to a remote location.

Figure 16 is an exploded partial elevational perspective view of a dispenser in accordance with the removable cassette embodiment of the present invention.

Figure 17 is a partial sectional view of the cassette of Figure 16, shown inserted into the dispenser.

Detailed Description of Preferred Embodiments

Referring to Figure 1, there is illustrated a dispensing kit 10 according to one embodiment of the present invention. The dispenser 10 comprises a main housing 11 for enclosing the reservoirs of blood proteins and working parts of the dispenser. Housing 11 may be formed from any of a variety of materials and by any of a variety of processes as will be appreciated by one of skill in the art. Preferably, housing 11 is produced in two halves in accordance with known thermoplastic forming techniques, and optimally comprises acrylic, styrene, or other rigid thermoplastic material that is susceptible to sterilization and will remain substantially inert in the environment of the blood proteins to be dispensed.

Housing 11 comprises a container region 12 having a plurality of container receiving areas 14, 16 (illustrated in Figure 2) adapted for removably receiving a plurality of containers 18, 20 (Figure 3) for containing the fibrinogen and thrombin blood proteins. Optimally, a thermal insulation material 22 such as styrofoam or other conveniently formable insulating material is provided to surround the container receiving areas 14, 16 to maintain the blood proteins at an appropriate preselected temperature as will be discussed.

The container receiving areas 14, 16 are optimally further provided with a cover 24 for preventing accidental removal of the containers 18, 20. Cover 24 is provided

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with a hinge 26 and a latch 28 for engaging stop 30 on the housing for preventing accidental removal of the cover 24. Other convenient hinge and latch arrangements can be readily envisioned by one of skill in the art. Providing
5 the housing 11 with a hinged cover 24 having a reversible latch 28 facilitates removal of empty containers 18, 20 and replacement with filled containers (not illustrated).

Referring to Figure 4, the container region 12 of the dispensing kit 10 may further be provided with a window 32
10 for visual observation of the amounts of blood proteins remaining in the containers 18, 20.

Adjacent the container region 12 of the dispensing kit 10 is a pump region 34 which houses the pumping means for extracting a quantity of blood protein from each of the
15 containers 18, 20 and delivering them to the application site. Thus, within the pump region 34 are contained a first pump 44 and a second pump 46. Pumps 44, 46 may be any of a variety of commercially available pumps, including, for example, a linear piston-type pump, as
20 illustrated in the embodiment shown in Figure 1, or a trigger sprayer-type pump, as illustrated in the embodiment of Figure 3.

The particular pumping mechanisms to be utilized in the present invention is not critical, provided they may be
25 conveniently ganged to a common actuator, and are susceptible to sterilization for entry into a sterile field. One example of a suitable trigger type sprayer is the Model TS-800 Trigger Sprayer from Calmar, Inc. of Watchung, N.J., which may conveniently be adjusted from a focused stream to
30 a spray.

In the event that linear piston-type pumps, as illustrated in Figure 1, are utilized, an actuator head 42 is additionally provided to form a unitary spray head 36 so that both the first and second pumps may be simultaneously
35 operated. Actuator 42 can be molded from any of the same rigid thermoplastic materials used to mold the housing 11. Effluent ducts 70 and 72 are preferably adjustable so that

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each of the spray patterns illustrated in Figures 11-13 may be achieved. The ability to adjust the droplet size and the cross sectional area of interaction between the two effluent streams provides important flexibility to the clinician. The speed of formation of the fibrin polymer, and perhaps its strength are enhanced by a spray which permits intimate intermingling of droplets of the two fluids close to or at the application site.

Optimally, actuator 42 is removably linked to each of pumps 44 and 46 so that the pumps may be individually operated at the option of the clinician. In applications where hemostasis is sought, an actuator 42 or its functional equivalent will likely be preferred. However, it may be desirable in some circumstances to individually dispense the fluids, such as one on each opposing surface of an incision. This will delay the rapid polymerization until the two surfaces are brought into contact.

Referring to Figure 5, there is provided an interface device 50 adapted for placing a standard blood bag 52 or other container provided with a piercable septum 56 in fluid communication with a container 18 suitable for use in the dispensing kit 10 of the present invention. The interface 50 is provided at one end with a cannula 54 adapted for piercing the septum 56 contained on a standard blood bag 52. The interface 50 is further provided with a valve 58 which may be a stopcock, a pinch valve or any of a variety of known valving mechanisms for controlling the flow from the blood bag 52 into the container 18. The interface 50 is further provided with a hollow needle 60 in fluid communication by way of valve 58 with cannula 54, and adapted for piercing the resealable septum of a standard Vacutainer® or other vacuum bottle.

Evacuated sterile container 18 is typically provided with a piercable septum 19 for maintaining a vacuum therein. In use, the interface 50 will be placed in fluid communication with the fluid contents of blood bag 52 by way of cannula 54. With valve 58 in the closed position,

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needle 60 on interface 50 is inserted through resealable septum 19 on container 18. When valve 58 is open, fluid contents from blood bag 52 are drawn into the interior of container 18 due to the vacuum contained therein. When
5 sufficient contents have been transferred into the container 18, valve 58 is closed and container 18 is removed from needle 60. Container 18 is then ready for insertion into one of container receiving areas 14, 16 on dispensing kit 10.

10 As container 18 is inserted into one of container receiving areas 14 or 16, the resealable septum 19 is first pierced by one of aspirating needles 62, 64. Aspirating needles 62, 64 are in fluid communication by way of pumps 44, 46 with effluent ducts 70, 72 on spray head 36.
15 Aspirating needles 62, 64 are disposed such that when containers 18, 20 are fully inserted into the container receiving areas 14, 16, the needles will reach substantially to the bottom of the interior of the containers.

20 As the container 18 is further inserted into the container receiving area 14, resealable septum 19 will be pierced by a second vent needle 66. A vent needle 66, 68 is desirable to equalize pressure inside containers 18, 20, as fluid contents are drawn out through needle 62 or 64.

25 As an alternative to Vacutainer® containers 18 and 20, nonvacuum bottles could also be used in the dispensing kit 10 of the present invention. Sample bottles can be used which have a threaded neck such as to receive a cap, or which are provided with a luer or other mechanical interfit
30 connection. The innermost extent of container receiving areas 14, 16 would be provided with a corresponding structure for engaging the luer or other interfit structure on the sample bottle. In this embodiment, it would be unnecessary for the aspirating needles 62, 64 to have
35 sharpened ends.

In accordance with a further embodiment of the present invention, there is provided a modified dispensing kit

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adapted to receive the containers of fibrinogen and thrombin within a removable cassette. Referring to Figure 16, there is illustrated in exploded form a dispenser 118 and cassette 122, in accordance with the present invention. As illustrated therein, housing 120 of dispenser 118 is adapted to slidably receive the upper end 126 of cassette 122. Cassette 122 may be produced in accordance with any of a variety of known methods, such as thermoplastic forming techniques, and optimally comprises acrylic, styrene or other rigid thermoplastic material. Cassette 122 may conveniently be formed in the same manner and of the same materials as the dispenser 10 described previously. In addition, cassette 122 is provided with insulation 132 for surrounding containers 123 and 124 as has been discussed.

The lower end 128 of cassette 122 is provided with a door 130 for providing access to the interior of the cassette 122. Door 130 is illustrated as a slidable panel, however, any of a variety of doors may be used as will be appreciated by one of skill in the art. Opening door 130 permits access to the interior of cassette 122 for insertion of containers 123 and 124, illustrated in phantom in Figure 16. The upper end 126 of cassette 122 is provided with openings 125 and 127 to expose the containers 123 and 124 for providing fluid communication with the pumps, as will become apparent.

Cassette 122 is further provided with a pair of opposing surfaces 138 for providing friction so that the cassette may be conveniently gripped, such as between the thumb and forefinger, and inserted or withdrawn with respect to housing 120. Housing 120 is provided with a corresponding pair of openings 140 which expose the surfaces 138 when the cassette 122 is fully inserted into housing 120. Any of a variety of additional structures for facilitating removal of cassette 122 may be used, as will be apparent to one of skill in the art.

Referring to Figure 17, there is disclosed a partial

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elevational cross-sectional view of a cassette embodiment of the dispenser of the present invention, illustrating the cassette 122 inserted into the housing 120. As seen therein, cassette 122 is dimensioned so that its exterior wall permits a snug fit with the interior of housing 120 for retaining the cassette therein. In addition, static friction will exist between each of needles 134 and 136; and the piercable septums on containers 123 and 124, respectively, which are arranged in a similar manner as the needles and septums or other closures previously discussed. Any of a variety of structures well known to one of skill in the art may additionally be utilized for retaining cassette 122, such as releasably interlocking ridges and grooves on the interior surface of housing 120 and exterior surface of cassette 122 as will be appreciated by one of skill in the art.

As cassette 122 is inserted into housing 120, needles 134 and 136 pierce the corresponding closure members on containers 123 and 124, as has been previously discussed. For simplification purposes, vent needles, which have been discussed in connection with previous figures, have been deleted from Figure 17.

The removable cassette embodiment accrues the important advantage in the operating room of providing a convenient bridge across the boundary of the sterile field. The dispensing kit 118, including the exterior of cassette 122 must be maintained sterile throughout a procedure in the operating room. However, the exterior of the containers themselves will not be sterile. The embodiment of Figures 16 and 17 enables the sterile nurse or physician to hold cassette 27 while the circulating nurse inserts filled containers 123 and 124 into cassette 122. The cassette 122 can then be fit onto housing 120 without compromising the integrity of the sterile field.

In operation, a nurse or other attendant will typically reconstitute lyophilized thrombin from bovine or other source by mixing with water or isotonic saline

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diluents in accordance with known techniques. Lyophilized thrombin may be commercially obtained, for example, from the Parke-Davis division of Warner-Lambert Co. (Morris Plains, NJ) which markets bovine thrombin in 5, 10 and 20 ml vials under the name Thrombostat®.

The reconstituted thrombin is then warmed to and maintained at approximately 37° celsius in the container 18 or 20 for installation in the dispenser 10. The fibrinogen is warmed from its refrigerated storage temperature to approximately 37° celsius and loaded into a blood bag if not so stored, or other suitable storage apparatus to be suspended from an IV pole in a location convenient to the operating table or other procedure site.

An interface device 50 is provided for the fibrinogen bag, and a fibrinogen container 18 is then filled as previously described and inserted into the dispensing kit 10 along with a bottle of reconstituted thrombin. Fibrinogen is typically available on an autologous basis or from a single donor source of plasma, which will be fractionated by a blood bank to produce a fibrinogen concentrate. Although fibrinogen extracts from multiple sources are commercially available in some countries, they are not yet in the United States due to certain prescreening difficulties known to those of skill in the art.

In the course of the surgical procedure, if additional fibrinogen and thrombin are necessary, additional vacuum bottles or other container 18 can be conveniently loaded from blood bag 52 or reconstituted and inserted into the dispensing kit 10 as needed. The ability to refill the dispensing kit 10 from a single source of each of the fibrinogen and thrombin accrues important advantages, particularly under circumstances when single source or autologous fibrinogen is being used.

As previously discussed, the fibrin polymer formed from the interaction of fibrinogen and thrombin may be advantageously applied in a variety of clinical situations.

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In addition to use as a tissue adhesive either during surgery or as a temporary emergency measure, and to the use of the fibrin polymer as a hemostatic agent, another important application relates to the implantation of vascular prostheses. Synthetic tubular grafts available for use in vascular surgery have been surgically implanted in hundreds of thousands of individuals over the course of the last decade. One such graft is comprised of rapidly stretched highly crystalline polytetrafluoroethylene, and is currently marketed by W. L. Gore & Associates, Inc.

One difficulty associated with the implantation of woven or stretched vascular prostheses is the occurrence of leakage when initially sutured into the vascular system. Leakage is an inherent result of the desirability of having a porous structure to permit interweaving of the tissue, and can particularly be a problem if the patient has been anticoagulated through the use of such agents as heparin or other anticoagulants utilized in the operating room. Hence, it is advantageous to be able to pretreat the porous vascular graft prior to implantation to prevent leakage therethrough during the period of time required for the tissue growth to render the implant fluid-tight.

Thus, there is additionally provided in accordance with the present invention a radial spray head 74 for uniformly dispensing each of two solutions containing fibrinogen and thrombin, respectively, onto the interior surface of a tubular sleeve 76, as illustrated in Figure 6. The tubular sleeve 76, illustrated in a cutaway view, may be any of a variety of commercially available vascular prostheses as previously discussed. The radial spray head 74 comprises a first spray element 82 and a second spray element 84 in fluid communication by way of a first influent passage 78 and a second influent passage 80, respectively, with a source of the fibrinogen and thrombin containing solutions. The radial spray head 74 may be conveniently connected by way of a delivery conduit 110, such as that illustrated in Figure 15, to the dispensing

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kit 10 of the present invention.

The radial spray head 74 is comprised of two spray elements 82, 84, which may be conveniently produced such as by a molding process to be identical to one another in configuration. For example, in Figure 7 there is illustrated a single spray element 82 which arbitrarily has been designated as the first spray element 82 corresponding to Figure 6. In the spray element 82, the influent passage 78 is in fluid communication with a plurality of effluent ducts 86 directed radially outwardly around the periphery of the spray element 82. Although the spray element 82 is illustrated as being substantially cylindrical in configuration, that is merely a preferred configuration due to the likelihood that most tubular vascular grafts will be of substantially circular cross section. However, for particular applications, it may be desirable to have a radial spray head 74 having a non-cylindrical configuration depending upon the particular application.

As illustrated in each of Figures 6-8, the spray elements 82 and 84 are each provided with a passageway 88 which performs no function in the distal spray element 84. In the proximal spray element 82, however, the passageway 88 permits stacking of the proximal and distal spray elements on top of one another by receiving the second influent passage 80 therethrough. This design is selected for manufacturing convenience and, of course, other configurations could readily be devised.

Referring to the partial cross-sectional view in Figure 8, it is seen that each of the influent passageways is respectively in fluid communication with the corresponding spray element. In a relatively large embodiment of the spray head 74, which may be desirable, for example, in applications utilizing larger arterial grafts, there may exist an undesirable dead space within the interior chamber 83 of element 82 so that a substantial quantity of fluid must be introduced into the spray element 82 before that element will express fluid uniformly around

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its periphery. Thus, it may be desirable to reduce the interior volume of the chamber 83. This can be accomplished, for example, by molding the spray element 82 such that only a relatively small annular flow passageway remains around the periphery of the spray element 82, for providing fluid communication between influent passage 78 and each of the effluent ducts 80.

The size and spacing of the effluent ducts 86 illustrated in Figures 6-8 is for illustrative purposes only, and a wide variety of duct patterns and spacings could be utilized in accomplishing the objectives of the present invention. In an alternate embodiment, spray element 82 could be comprised of a top and a bottom half which form therebetween a single annular effluent duct extending radially all the way around the periphery of the spray element 82. The efficacy of a single annular groove embodiment would be dependent upon manufacturing tolerances, and the use of a plurality of drilled or molded effluent ducts 86 may therefore be superior.

In use, the radial spray head 74 may be placed in fluid communication with the dispensing kit 10 or other source of fluids by way of a delivery conduit 110 such as that illustrated in Figure 15. Prior to implantation of the vascular graft, the radial spray head 74 may then be positioned within the vascular graft and drawn therethrough as fluid is expressed from the dispensing kit 10. In this manner, uniform quantities of the fibrinogen and thrombin solutions will be dispersed along the interior surface of the tubular sleeve 76, thereby forming a fluid-impermeable layer thereon.

As an alternative to or in addition to the interior spray head 74, there has further been provided in accordance with the present invention a spray element 90 for uniformly dispensing a plurality of fluids along the exterior of a tubular sleeve 76 (see Figure 9). The radial spray head 90 comprises a first spray element 96 in communication with a source of fluid by means of first

influent passage 92, and a second spray element 98 in communication with a source of fluid by way of second
influent passage 94. Influent passages 92, 94 may be
similarly placed in fluid communication with a source of
5 fluids such as dispensing kit 10 by way of a delivery
conduit 110. The first and second spray elements 96, 98
may be identical in configuration, and a single spray
element 96 is illustrated in Figure 10. Referring to
Figure 10, the first influent passage 92 is in fluid
10 communication by way of an annular chamber (not
illustrated) in spray element 96 with a plurality of
effluent ports 100 dispersed radially around the interior
of the torus-shaped element 96. The cylindrical opening
through spray head 96 has a diameter to facilitate
15 insertion of tubular sleeve 76 therethrough. As with the
spacing of the effluent ducts 86 on the spray head 74
illustrated in Figure 6, the frequency and configuration of
the effluent ports 100 can vary considerably, however, it is
contemplated that about ten or more effluent ports per inch
20 of arc length along the interior cylindrical surface 101 of
spray element 96 will be used.

Each element 96, 98 of the spray head 90 is
illustrated with the influent passage disposed at a
perpendicular to the plane of the spray element. This
25 configuration may conveniently be accomplished by
displacing the axis of effluent passage 92 radially
outwardly from the exterior diameter of spray element 96.
In this manner, identical spray elements can be stacked one
on top of another without interference from influent
30 passageways.

Of course, the configuration can be modified in a
variety of ways and still accrue the advantages of the
present invention. For example, the axis of effluent
passageway 92 could coincide with the annular chamber
35 within spray element 96. In that embodiment, the spray
element 96 would need to be provided with an opening
conceptually equivalent to passageway 88, illustrated in

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Figures 6-8, to permit stacking of the spray heads 96, 98. Alternatively, it may be desirable for the influent passageways 92, 94 to extend radially outwardly from the spray elements 96, 98 such that they extend at a perpendicular from the tubular sleeve 76 when in use. In this embodiment, the influent passageways could be integral with a handle for convenient operation of the spray head.

Use of the interior spray head 74 and exterior spray head 90 is advantageously accomplished through the use of a delivery conduit 110, illustrated in Figure 15. Delivery conduit 110 is a dual channel conduit comprised of a flexible, resilient material which is substantially inert in the presence of the fibrinogen and thrombin solutions to be conducted therethrough. Distal end 112 can be secured in fluid communication with a spray head, such as that illustrated in Figures 6 or 9, by friction fit with the influent passageways thereof. Alternatively, any of a variety of other known connection means can be employed.

Similarly, proximal end 114 of delivery conduit 110 is advantageously dimensioned to enable a friction fit to provide fluid communication with ducts 70 and 72 on spray head 36. Alternatively, proximal end 114 and spray head 36 can be provided with positive interlocking structures such as a luer connection or other known lock.

The delivery conduit 110 can be provided in a variety of different lengths, depending upon the intended use. Typically, vascular grafts for human implantation may be anywhere from about 10 cm to as long as 30 cm in length, or longer, and conduits of that length or less are currently envisioned. In another application of the present invention, a flexible delivery conduit 110 is provided having sufficient length to extend through the working channel in any of a variety of instruments such as an endoscope, arthroscope, or gastroscope which permits visualization at a remote site. This structure permits visualization and release of the fibrinogen and thrombin solutions at a remote spot in any of a variety of body

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cavities where hemostasis or an adhesive may be desired, such as a treatment for ulcerations in the stomach or gastrointestinal tract, or as a postoperative measure following such procedures as a tonsillectomy, adenoidectomy or more remote surgery. In a further embodiment, delivery conduit 108 (Figure 14) is typically shorter than conduit 110, and is rigid for precisely controlled delivery at a more accessible specified point.

Although this invention has been described in terms of certain preferred embodiments, other embodiments that are apparent to those of ordinary skill in the art are also within the scope of this invention. Accordingly, the scope of the invention is intended to be defined only by reference to the appended claims.

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I CLAIM:

1. A dispenser for separately dispensing each of two biological fluids contained therein for intermixing at a remote site to produce hemostasis or a tissue adhesive,
5 comprising:

a housing;
a first and a second pump in the housing;
a first hollow aspirating needle in the housing,
in fluid communication with the first pump;
10 a second hollow aspirating needle in the housing,
in fluid communication with the second pump; and
a chamber in the housing for removably receiving
a first and a second reservoir of biological fluids to
be dispensed;

15 wherein each of the first and second aspirating
needles extends into the chamber for fluid
communication with each of the first and second
reservoirs of biological fluid, respectively.

2. A dispenser as in Claim 1, wherein each of the
20 first and second aspirating needles terminates in a sharp
point, for piercing a septum on the reservoir of biological
fluid.

3. A dispenser as in Claim 2, wherein the reservoir
comprises a vacuum bottle having a piercable, resealable
25 septum.

4. A dispenser as in Claim 1, further comprising a
first and second hollow vent needles in the housing,
extending into the chamber substantially parallel to the
first and second aspirating needles, respectively, for
30 communicating with the interior of the reservoirs.

5. A dispenser as in Claim 1, further comprising a
common actuator for simultaneously actuating each of said
first and second pumps.

6. A dispenser as in Claim 5, wherein the actuator
35 is removable secured to the pumps to permit removal thereof
for independent operation of the pumps.

7. A dispenser as in Claim 1, further comprising a

cover on the housing, for enclosing the chamber to prevent accidental removal of the first and second reservoirs of biological fluids.

8. A dispenser as in Claim 7, wherein the cover is
5 pivotally attached to the housing.

9. A dispenser as in Claim 8, wherein the cover further comprises a latch for securing the cover in the closed position.

10. A dispenser as in Claim 1, further comprising a
10 layer of thermal insulating material disposed on the inside surface of the chamber.

11. A dispenser as in Claim 1, further comprising a
15 spray head having a first and a second effluent ducts, in fluid communication with the first and second pumps, respectively, for directing a first and second stream of biological fluid for intermingling at a target site.

12. A dispenser as in Claim 11, wherein each of the
20 first and second effluent ducts is adjustable from a first position for producing a steady stream of fluid, to a second position for producing an aerosol mist.

13. A dispenser as in Claim 11, wherein the effluent
25 ducts are adjustable from a first position wherein the effluent streams converge at a first distance from the spray head, to a second position wherein the effluent streams converge at a second distance from the spray head.

14. A dispenser as in Claim 7, wherein the cover is
30 removably secured to the housing, and the cover comprises a cavity adapted to receive at least one of the reservoirs so that the cover may be loaded with the reservoirs and then installed as a unit into the housing.

15. An interface device for placing a vacuum bottle
having a resealable septum thereon in valved fluid communication with a source of fluid, comprising:

35 a housing having a flow channel therethrough;
a valve in the flow channel;
a hollow needle in communication with the flow channel on a first side of the valve; and

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a hollow cannula in communication with the channel on a second side of the valve;

wherein the needle is adapted for piercing the septum on a vacuum bottle.

5 16. A radial spray head for dispensing each of a plurality of fluids onto the surface of a tubular sleeve, comprising:

10 a first spray element, comprising an influent passageway in fluid communication with a plurality of effluent ducts disposed about the periphery of the first spray element; and

15 a second spray element comprising a second influent duct in fluid communication with each of a plurality of effluent ducts disposed about the periphery of the second spray element.

20 17. A radial spray head as in Claim 16, for uniformly distributing each of two fluids along the interior surface of a tubular vascular graft, wherein the plurality of effluent ducts are disposed about the exterior periphery of the first and second spray elements for directing a spray in a radially outward direction.

25 18. A radial spray head as in Claim 16, for uniformly distributing each of two fluids along the exterior surface of a tubular vascular graft, wherein the plurality of effluent ducts are disposed about the interior periphery of a tubular aperture through the first and second spray elements, for directing a spray in a radially inward direction.

30 19. A dispenser for separately dispensing each of two biological fluids contained therein for intermixing at a site outside of the dispenser to produce hemostasis or a tissue adhesive, comprising:

35 a dispenser housing having a chamber for removably receiving a cassette for holding a first and a second reservoir of biological fluids to be dispensed;

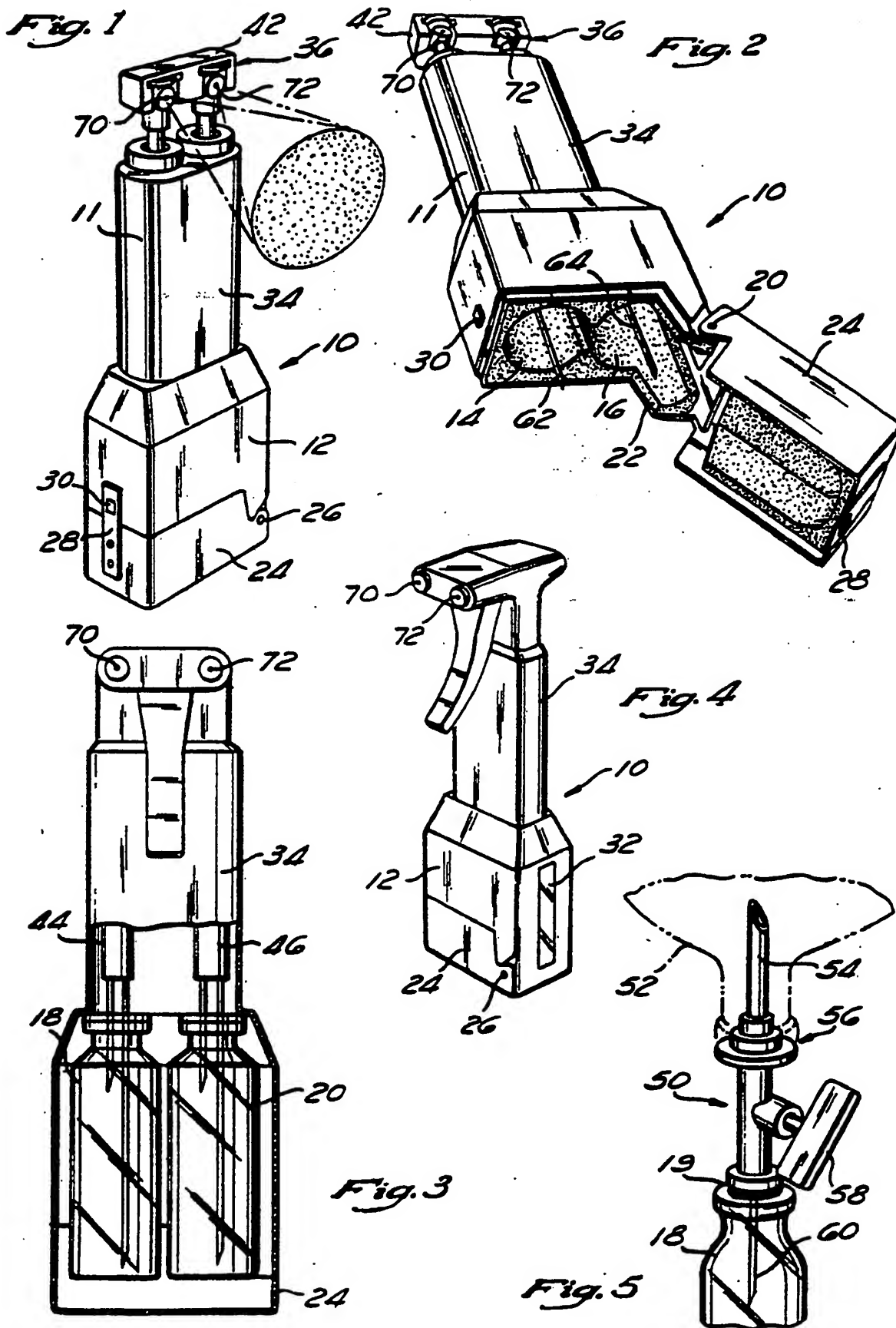
a first and a second pump in the housing;

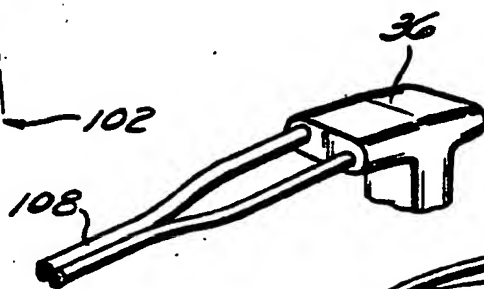
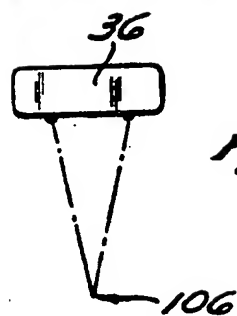
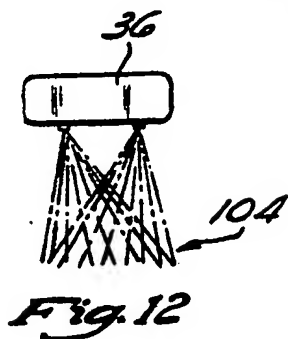
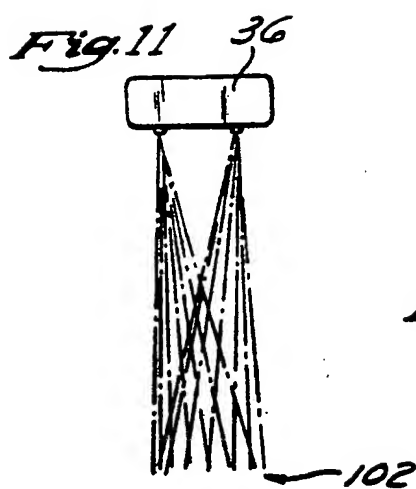
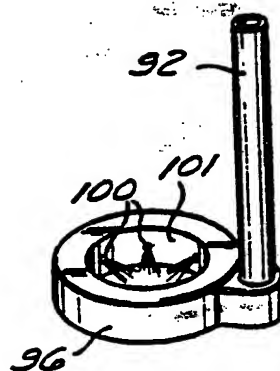
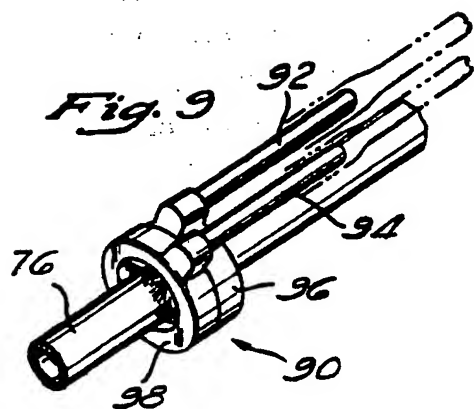
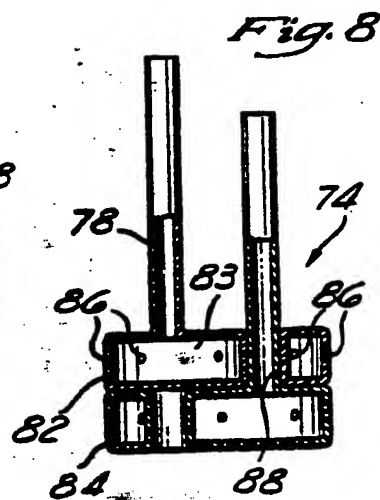
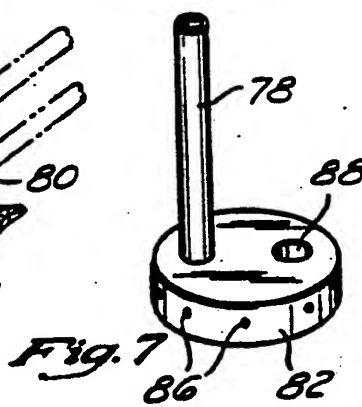
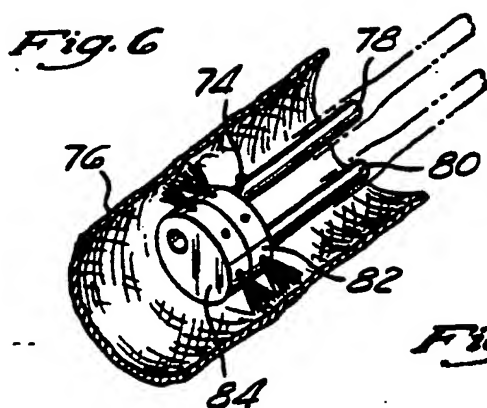
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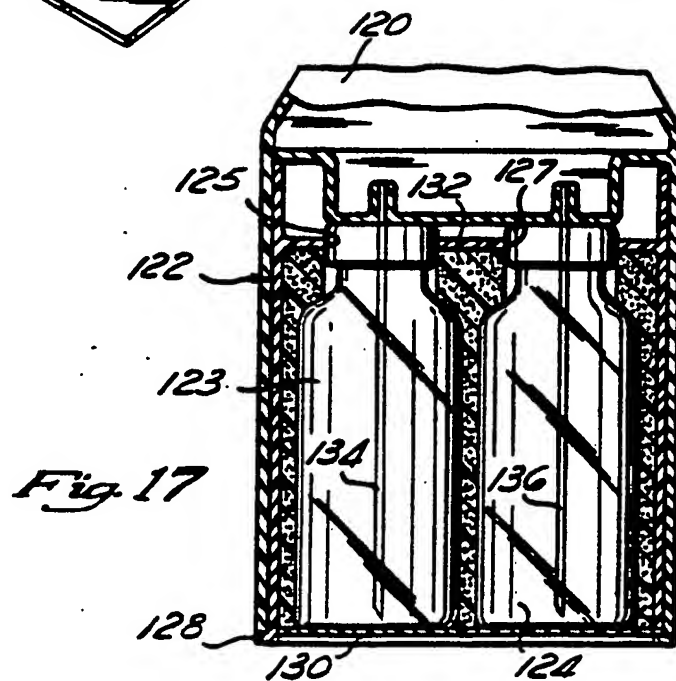
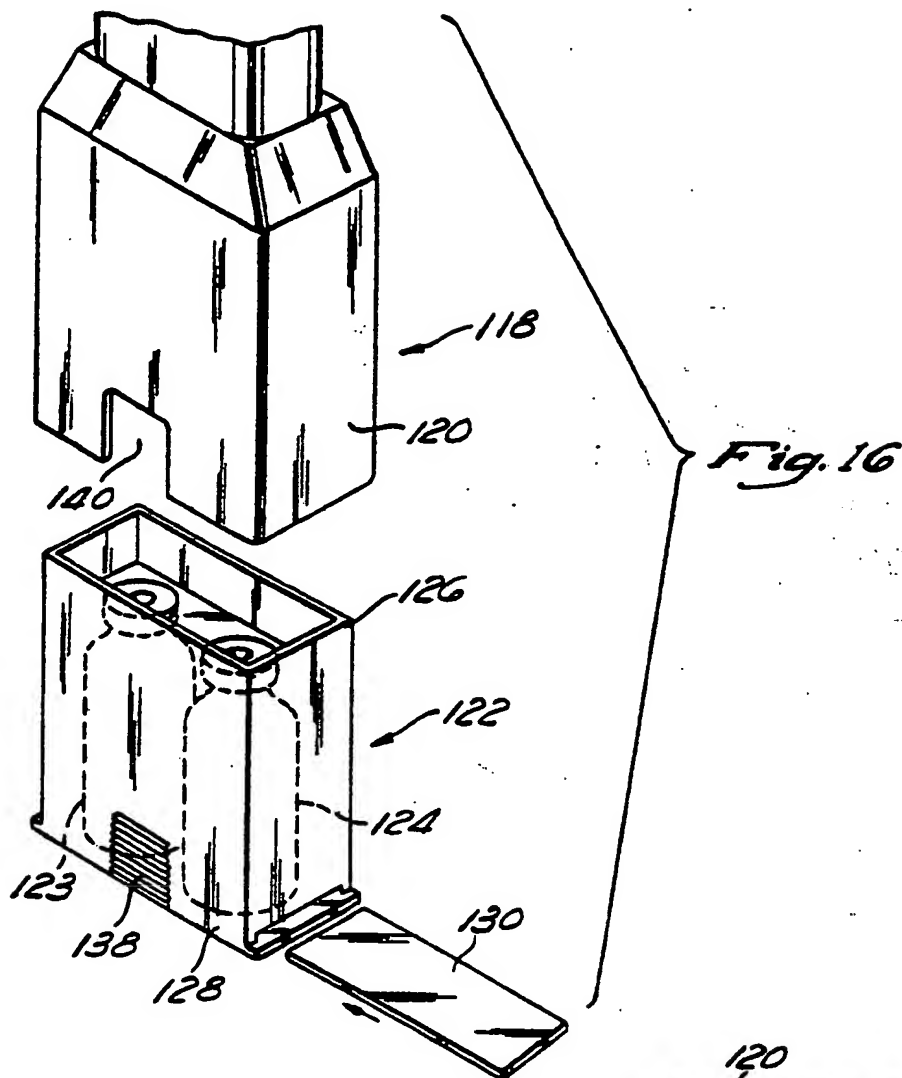
a first hollow aspirating needle in the housing,
in fluid communication with the first pump and
extending into the chamber; and

5 a second hollow aspirating needle in the housing,
in fluid communication with the second pump and
extending into the chamber;

10 wherein the cassette comprises a housing for
containing a reservoir of each of the two biological
fluids, and means for releasably engaging the
dispenser housing so that the first and second pumps
are placed in fluid communication with the first and
second reservoirs, respectively, when the cassette is
engaged with the dispenser housing.







INTERNATIONAL SEARCH REPORT

International Application No. PCT/US89/03468

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC		
IPC (4) A61M 5/00		
US CL 604/191		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
	604/82, 83, 88, 191, 201	
US	222/135, 137, 81, 94, 255	
Documentation Searched other than Minimum Documentation to the extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	US,A, 4,735,616 (EIBL ET AL), 05 APRIL 1988.	1
Y	US,A, 4,631,055 (REDL ET AL) (See Abstract), 23 DECEMBER 1986.	1-19
Y	US,A, 3,767,085 (CANNON ET AL), 23 OCTOBER 1973. See cols. 7, 8, lines 25-64, lines 1-11.	1-19
A	US,A, 4,040,420 (SPEER), 09 AUGUST 1977.	
<p>¹⁰ Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"G" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search		Date of Mailing of this International Search Report
30 October 1989		21 NOV 1989
International Searching Authority		Signature of Authorized Officer
		<i>[Signature]</i>